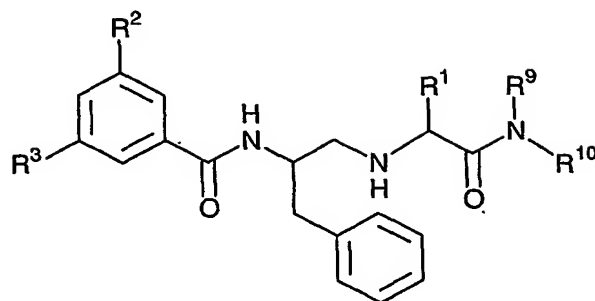


WHAT IS CLAIMED IS:

1. A compound of the formula I:



I

wherein:

R¹ is selected from the group consisting of:

- (1) C₁₋₆alkyl, unsubstituted or substituted with -OR⁵ or -S(O)₂-C₁₋₆alkyl,
- (2) hydrogen,
- (3) phenyl, and
- (4) benzyl;

R² is selected from the group consisting of:

- (1) hydrogen,
- (2) R⁴-S(O)_p-,

wherein R⁴ is independently selected from the group consisting of:

- (a) -C₁₋₆alkyl, which is unsubstituted or substituted with 1-6 fluoro,
- (b) phenyl, and
- (c) benzyl,

- (3) R⁴-S(O)_pN(R⁵)-,

wherein R⁵ is independently selected from the group consisting of:

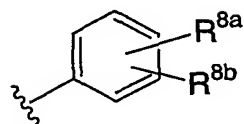
- (a) hydrogen,
- (b) -C₁₋₆alkyl, which is unsubstituted or substituted with 1-6 fluoro,
- (c) -C₃₋₆cycloalkyl which is unsubstituted or substituted with methyl,
- (d) phenyl, which is unsubstituted or substituted with halo or methoxy, and
- (e) benzyl,

- (4) -CN,

(5) -C₁₋₆alkyl-CN,

(6) halogen,

(7)



5 wherein R^{8a} and R^{8b} are independently selected from the group consisting of:

(a) hydrogen,

(b) -CN,

(c) halo,

(d) -C₁₋₆alkyl,

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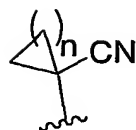
(e) -O-R⁵,

(f) -S-R⁵,

(g) -CO₂R⁵, and

(h) tetrazolyl,

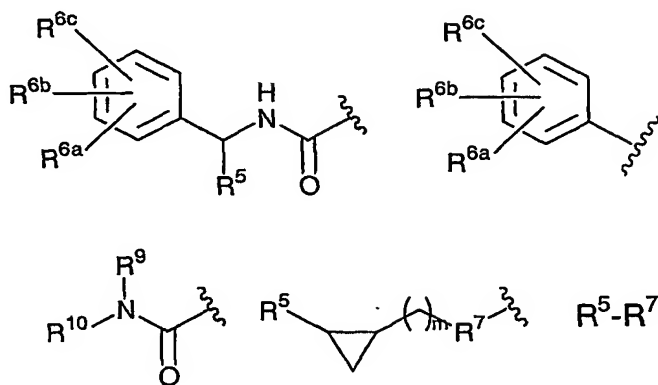
(8)



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wherein n is 1, 2, 3 or 4;

R³ is selected from the group consisting of:



;

20 R^{6a}, R^{6b}, and R^{6c} are independently selected from the group consisting of:

(1) hydrogen,

- (2) halogen,
- (3) -OR⁵,
- (4) -SR⁵, and
- (5) -C₁₋₆alkyl;

5

R⁷ is selected from the group consisting of a bond, -CH=CH-, -O-, -S-, and -NH-;

R⁹ and R¹⁰ are independently selected from the group consisting of:

- (1) hydrogen,
 - 10 (2) C₁₋₆alkyl, unsubstituted or substituted with -CN or 1-4 halo,
 - (3) -C₃₋₆cycloalkyl,
 - (4) phenyl, which is unsubstituted or substituted with halo or methoxy, and
 - (5) benzyl,
- or R⁹ and R¹⁰ may be joined together to form a pyrrolidine or piperidine ring which is
- 15 unsubstituted or substituted with benzyl, -OR⁵ or 1-4 halo;

m is independently 0, 1, or 2;

p is independently 0, 1, or 2,

and pharmaceutically acceptable salts thereof.

20

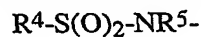
2. The compound of Claim 1 wherein R¹ is C₁₋₆alkyl.

3. The compound of Claim 1 wherein R¹ is methyl.

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4. The compound of Claim 1 wherein R¹ is ethyl.

5. The compound of Claim 1 wherein R² is:



and wherein R⁴ is selected from the group consisting of:

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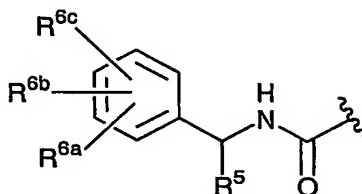
- (4) C₁₋₆alkyl,
- (5) phenyl, and
- (6) benzyl;

R⁵ is selected from the group consisting of:

- (5) C₁₋₆alkyl,

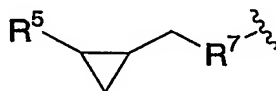
- (6) phenyl,
- (7) benzyl, and
- (8) hydrogen.

5 6. The compound of Claim 1 wherein R³ is:



and wherein R⁵ is methyl, R^{6a} is H or F, R^{6b} and R^{6c} are hydrogen.

7. The compound of Claim 1 wherein R³ is:

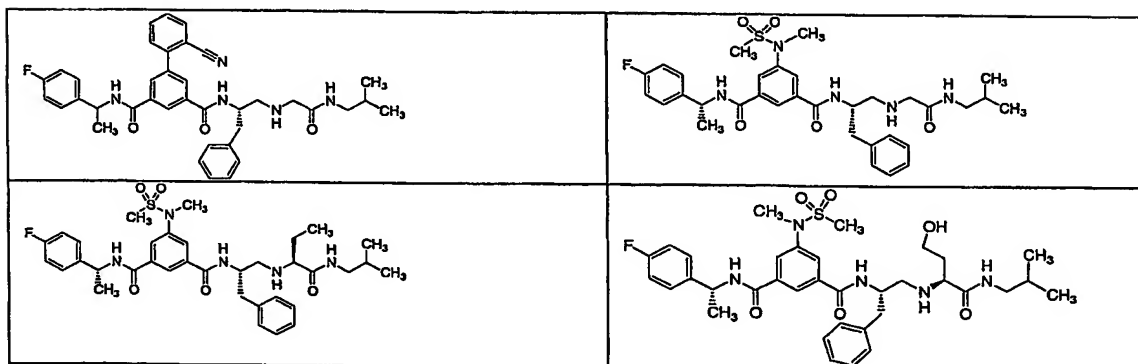


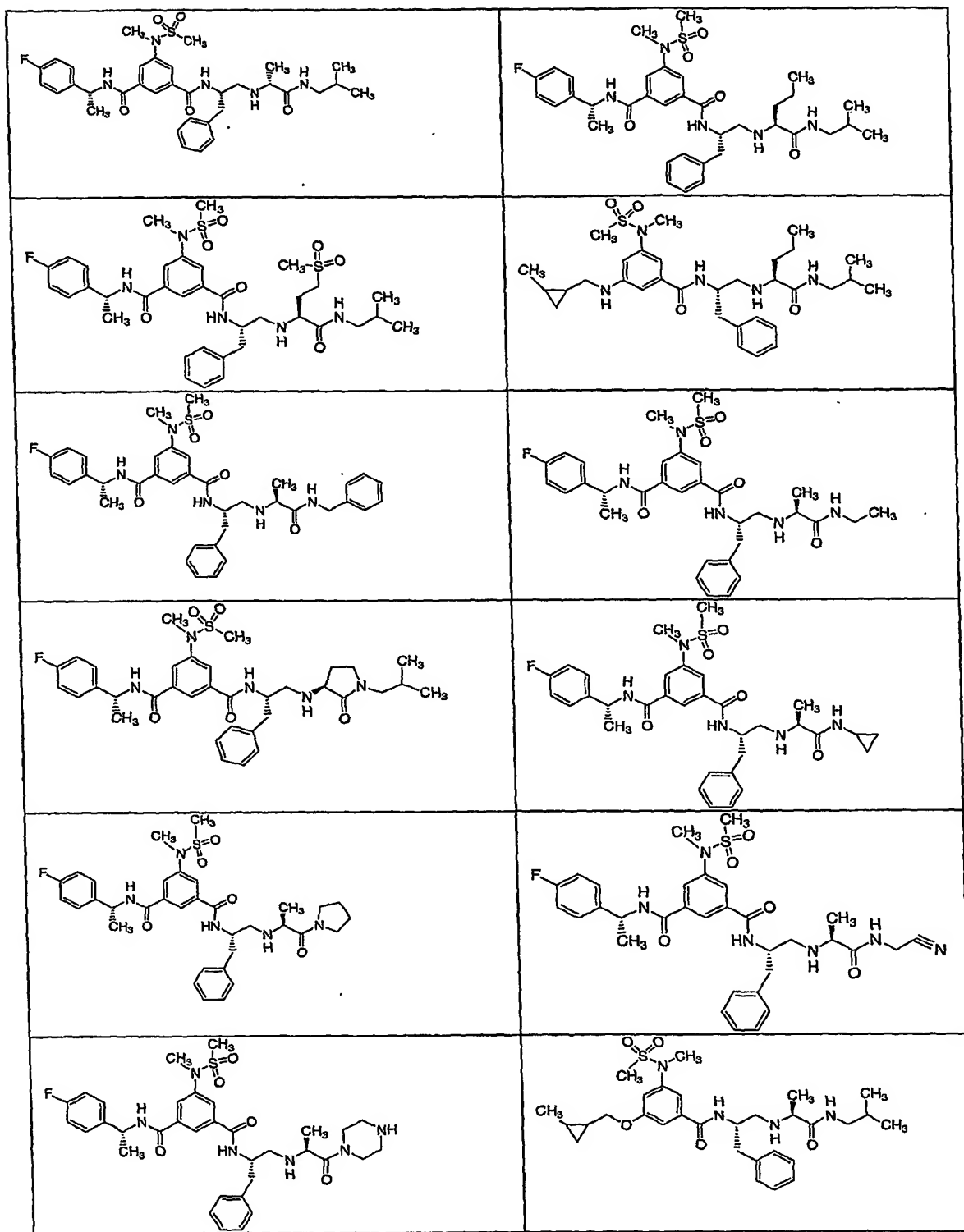
8. The compound of Claim 1 wherein R⁹ is hydrogen.

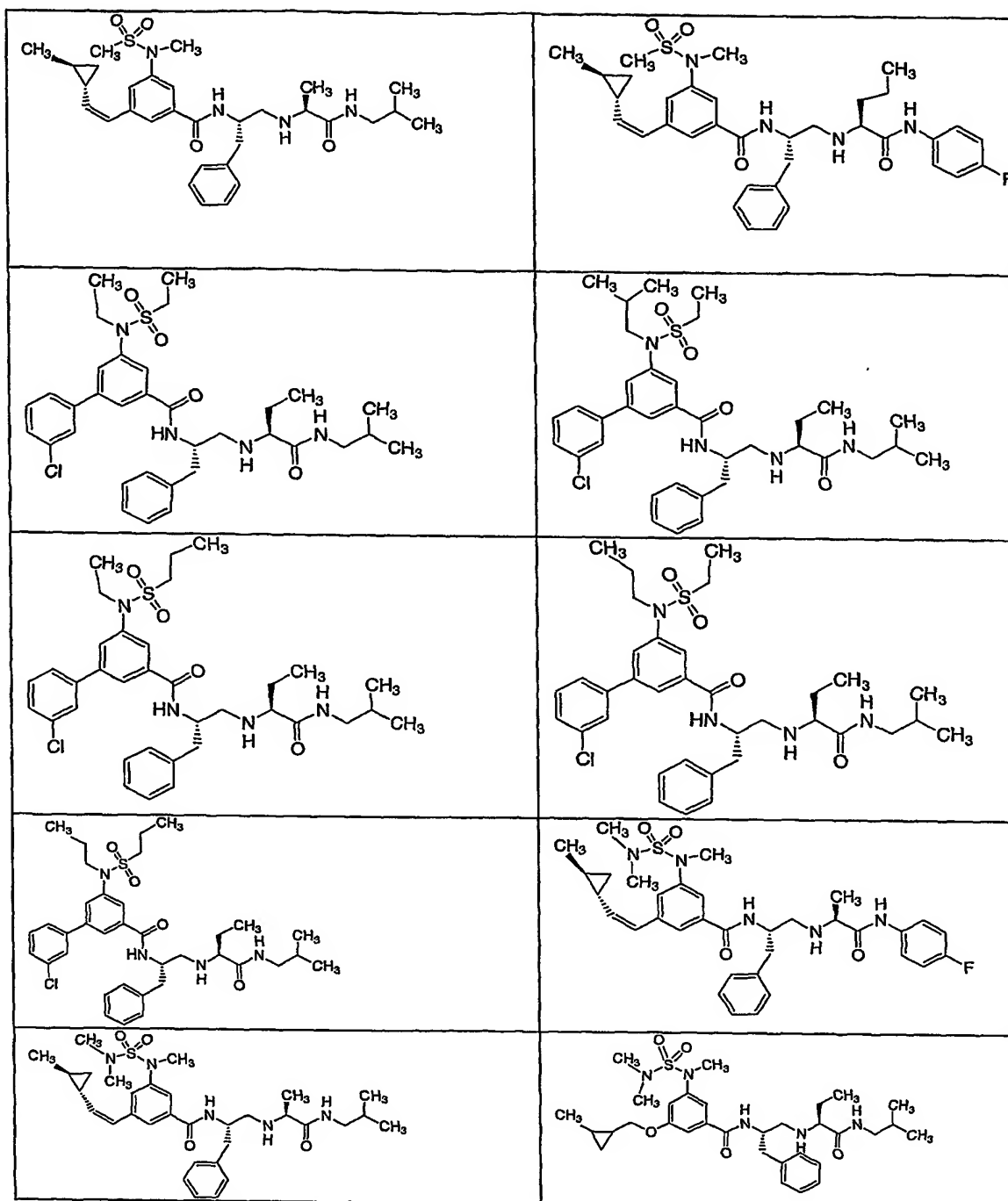
9. The compound of Claim 1 wherein R¹⁰ is C₁₋₆alkyl.

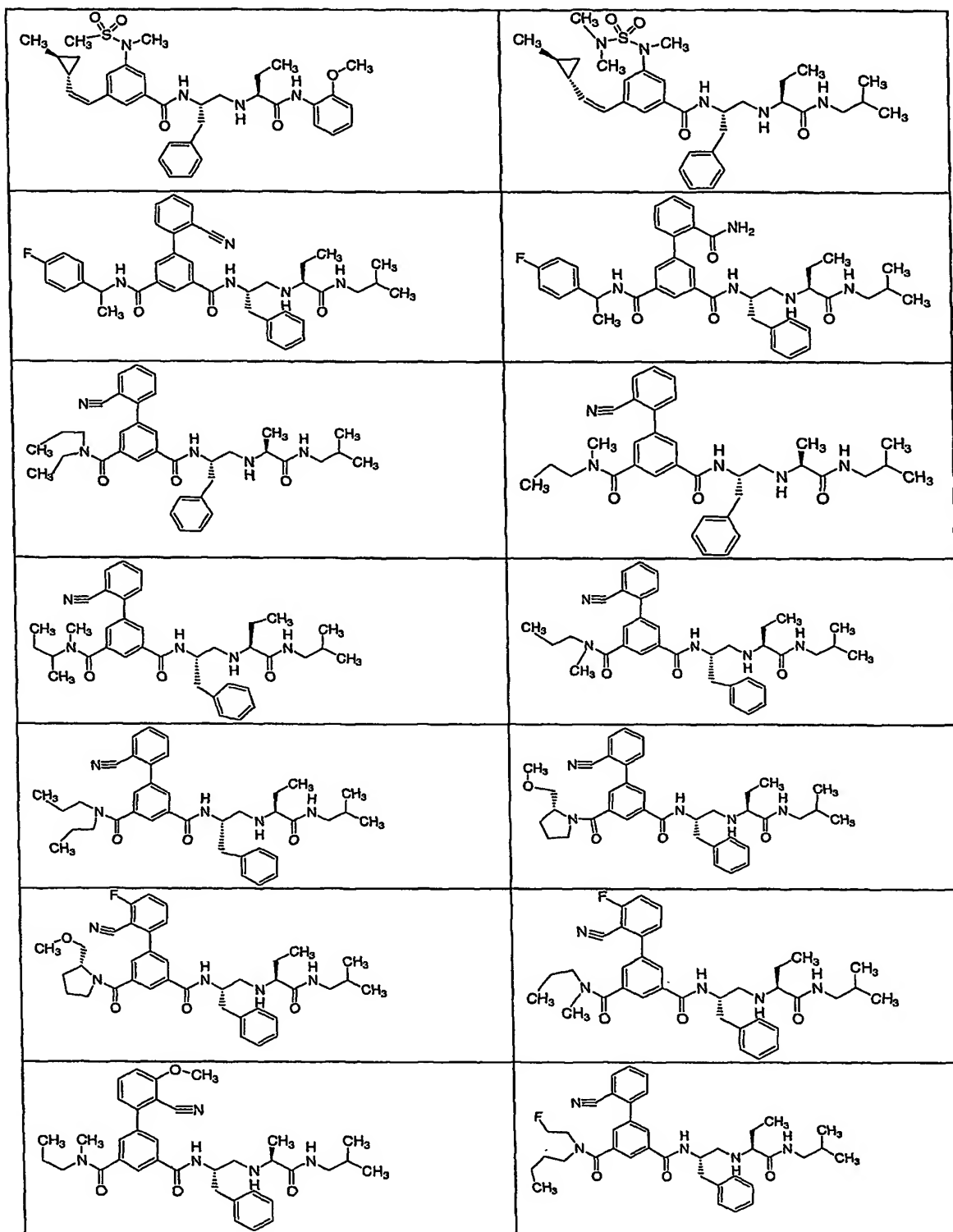
10. The compound of Claim 1 wherein R¹⁰ is iso-butyl.

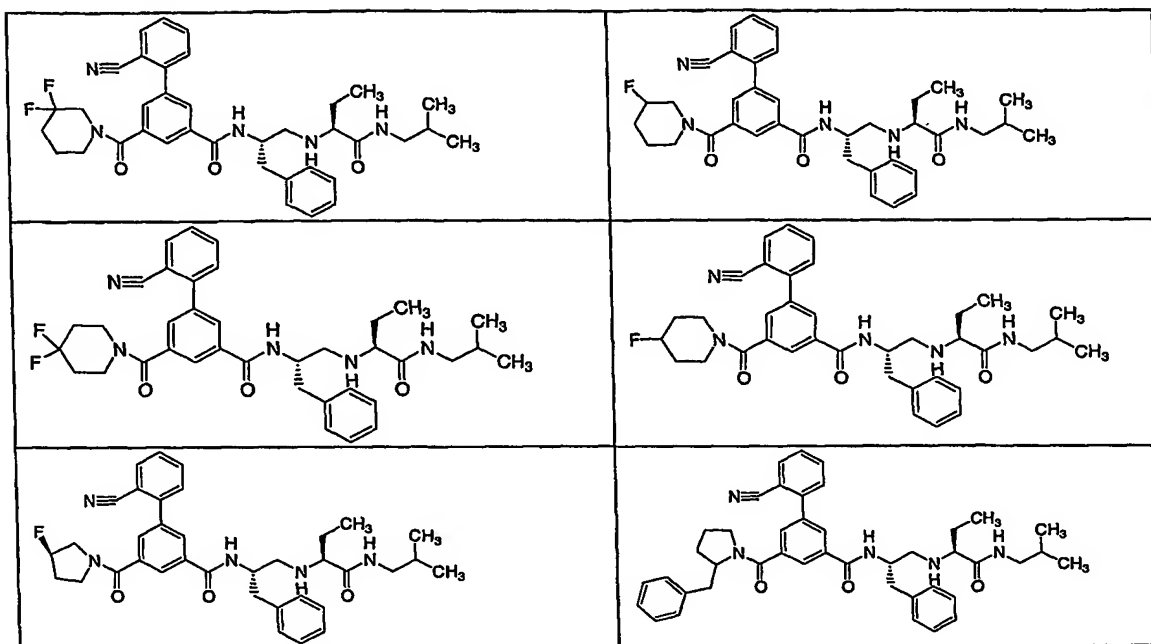
11. A compound which is selected from the group consisting of:











and pharmaceutically acceptable salts thereof.

12. A pharmaceutical composition comprising a therapeutically effective amount of the compound of Claim 1 and a pharmaceutically acceptable carrier.

13. A method for inhibition of β -secretase activity in a mammal which comprises administering to the mammal in need thereof a therapeutically effective amount of the compound of Claim 1 or a pharmaceutically acceptable salt thereof.

14. A method for treating Alzheimer's disease in a patient comprising the administration to the patient of a therapeutically effective amount of the compound of Claim 1 or a pharmaceutically acceptable salt thereof.

15. A method for preventing, controlling, ameliorating or reducing the risk of Alzheimer's disease in a patient comprising the administration to the patient of a therapeutically effective amount of the compound of Claim 1 or a pharmaceutically acceptable salt thereof.